IN THE CLAIMS:

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Please amend the claims as follows:

- 1. (Currently Amended) A method for prophylaxis or treatment of breast cancer in a mammalian patient comprising administering to said patient a therapeutically effective amount of one or more compound(s) selected from the group consisting of carbetocin and other longacting oxytocin analogues in a pharmaceutically acceptable carrier sufficient to inhibit initiation or growth of breast cancer in said patient.
 - 2. (Cancelled)
- 3. (Currently Amended) The method of claim 1, wherein said one or more oxytocin analogue(s) is/are carbetocin is administered to said patient by a mode of administration selected from intramuscular, intravenous, intranasal, intrapulmonary, subcutaneous, parenteral, oral, or transdermal delivery.
- 4. (Currently Amended) The method of claim 3, wherein said one or more oxytocin analogue(s) is/are carbetocin is administered to said patent intranasally.
- 5. (Currently Amended) The method of claim 3, wherein said one or more oxytocin analogue(s) is/are carbetocin is formulated in said carrier for intranasal or intrapulmonary administration.
- 6. (Currently Amended) The method of claim 5, wherein said one or more oxytocin analogue(s) is/are carbetocin is formulated in a powder or aqueous formulation for intranasal delivery.
- 7. (Currently Amended) The method of claim 6, wherein said one or more oxytocin analogue(s) is/are carbetocin is combined in an aqueous formulation with one or more excipients selected from the group consisting of nonoxynol-9, laureth-9, poloxamer-124, octoxynol-9, lauramide DEA, chlorobutanol, glycerol, citric acid, sodium acetate for intranasal delivery.
- 8. (Original) The method of claim 6, wherein said carbetocin is formulated with a nonionic surfactant and polysorbate-80 in an aqueous formulation for intranasal delivery.
- 9. (Currently Amended) The method of claim 1, wherein said one or more oxytocin analogue(s) is/are carbetocin is administered in a dose of at least one microgram.
- 10. (Currently Amended) The method of claim 1, wherein said one or more oxytocin analogue(s) is/are carbetocin is administered daily in an intranasal formulation.

- 11. (Currently Amended) The method of claim 1, further comprising administering tamoxifen and/or raloxifene to said patient in an amount sufficient to inhibit initiation or growth of estrogen-dependent breast cancer in said patient.
- 12. (Currently Amended) The method of claim 11, wherein said one or more oxytocin analogue(s) is/are carbetocin and said tamoxifen and/or raloxifene are administered simultaneously as a mixture.
- 13. (Currently Amended) A method of prophylaxis or treatment of a psychiatric obsessive compulsive disorder in a mammalian patient comprising administering to said patient a therapeutically effective amount of one or more compound(s) selected from the group consisting of carbetocin and other long-acting oxytocin analogues in a pharmaceutically acceptable carrier sufficient to alleviate an obsessive-compulsive behavior of said disorder in said patient.
 - 14. (Cancelled)

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- 15. (Cancelled)
- 16. (Currently Amended) The method of claim 13, wherein said one or more oxytocin analogue(s) is/are carbetocin is administered to said patient by a mode of administration selected from intramuscular, intravenous, intranasal, intrapulmonary, subcutaneous, parenteral, oral, and transdermal delivery.
- 17. (Currently Amended) The method of claim 16, wherein said one or more oxytocin analogue(s) is/are carbetocin is administered to said patient intranasally.
- 18. (Currently Amended) The method of claim 16, wherein said one or more oxytocin analogue(s) is/are carbetocin is formulated in said carrier for intranasal or intrapulmonary administration.
- 19. (Currently Amended) The method of claim 18, wherein said one or more oxytocin analogue(s) is/are carbetocin is formulated in a powder or aqueous formulation for intranasal delivery.
- 20. (Currently Amended) The method of claim 19, wherein said one or more oxytocin analogue(s) is/are carbetocin is combined in an aqueous formulation with one or more excipients selected from the group consisting of nonoxynol-9, laureth-9, poloxamer-124, octoxynol-9, lauramide DEA, chlorobutanol, glycerol, citric acid, sodium phosphate, methyl paraben, propyl paraben, sorbitol, sodium chloride, and/or sodium acetate for intranasal delivery.

- 21. (Original) The method of claim 19, wherein said carbetocin is formulated with a nonionic surfactant and polysorbate-80 in an aqueous formulation for intranasal delivery.
- 22. (Currently Amended) The method of claim 13, wherein said one or more oxytocin analogue(s) is/are carbetocin is administered daily in an intranasal formulation.
- 23. (Currently Amended) The method of claim 13, wherein said one or more oxytocin analogue(s) is/are carbetocin is administered daily in an intranasal formulation.
- 24. (Original) The method of claim 13, further comprising administering a selective serotonin reuptake inhibitor or serotonin reuptake inhibitor to said patient in an amount sufficient to alleviate an obsessive-compulsive behavior in said patient.
- 25. (Currently Amended) The method of claim 24, wherein said one or more oxytocin analogue(s) is/are carbetocin and said selective serotonin reuptake inhibitor are administered simultaneously as a mixture.
- 26. (Currently Amended) A pharmaceutical composition for prophylaxis or treatment of breast cancer in a mammalian patient comprising a therapeutically effective amount of one or more oxytocin analogue(s) selected from the group consisting of carbetocin and other longacting oxytoc in analogues in a pharmaceutically acceptable carrier, wherein said composition is sufficient to inhibit initiation or growth of breast cancer in said patent.
 - 27. (Cancelled)

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- 28. (Currently Amended) The pharmaceutical composition of claim 26, wherein said one or more oxytocin analogue(s) is/are carbetocin is formulated in said carrier for intranasal or intrapulmonary administration.
- 29. (Currently Amended) The pharmaceutical composition of claim 26, wherein said one or more oxytocin analogue(s) is/are carbetocin is formulated in a powder or aqueous formulation for intranasal delivery.
- 30. (Currently Amended) The pharmaceutical composition of claim 26, wherein said one or more oxytocin analogue(s) is/are carbetocin is combined in an aqueous formulation with one or more excipients selected from the group consisting of nonoxynol-9, laureth-9, poloxamer-124, oxtoxynol-9, lauramide DEA, chlorobutanol, glycerol, citric acid, sodium phosphate, methyl paraben, propyl paraben, sorbitol, sodium chloride, and/or sodium acetate for intranasal delivery.

- 31. (Original) The pharmaceutical composition of claim 26, prepared in a unit dosage form containing at least one microgram of said one or more oxytocin analogue(s).
- 32. (Currently Amended) The pharmaceutical composition of claim 26, further comprising tamoxifen and/or raloxifen in an amount sufficient to inhibit initiation or growth of estrogen-dependent breast cancer in said patient.
- 33. (Currently Amended) A medicament suspension or powder for nasal administration to treat or prevent breast cancer comprising carbetocin and a powder of one or more cation exchange resins and/or one or more absorbent resins.
- 34. (Currently Amended) A pharmaceutical composition for prophylaxis or treatment of a psychiatric an obsessive compulsive disorder in a mammalian patient comprising a therapeutically effective amount of one or more oxytocin analogue(s) selected from the group consisting of carbetocin and other long-acting oxytocin analogues in a pharmaceutically acceptable carrier, wherein said composition is sufficient to alleviate at least one symptom of said psychiatric obsessive compulsive disorder in said patient.
 - 35. (Cancelled)
- 36. (Currently Amended) The pharmaceutical composition of claim 34, wherein said one or more oxytocin analogue(s) is/are carbetocin is formulated in said carrier for intranasal or intrapulmonary administration.
- 37. (Currently Amended) The pharmaceutical composition of claim 34, wherein said one or more oxytocin analogue(s) is/are carbetocin is formulated in a powder or aqueous formulation for intranasal delivery.
- 38. (Original) The pharmaceutical composition of claim 34, further comprising a selective serotonin reuptake inhibitor or serotonin reuptake inhibitor.
- 39. (Currently Amended, Previously Added) A method for treatment of breast cancer in a mammalian patient comprising administering to said patient a therapeutically effective amount of one or more compound(s) selected from the group consisting of carbetocin and other long acting oxytocin analogues in a pharmaceutically acceptable carrier sufficient to inhibit growth of breast cancer in said patient.
 - 40. (Cancelled)
- 41. (Currently Amended) The method of claim 39, wherein said one or more oxytocin analogue(s) is/are carbetocin is administered to said patient by a mode of administration

selected from intramuscular, intravenous, intranasal, intrapulmonary, subcutaneous, parenteral, oral, or transdermal delivery.

- 42. (Currently Amended) The method of claim 41, wherein said one or more oxytocin analogue(s) is/are carbetocin is administered to said patient intranasally.
- 43. (Currently Amended) The method of claim 41, wherein said one or more oxytocin analogue(s) is/are formulated in said carrier for intranasal or intrapulmonary administration.
- 44. (Currently Amended) The method of claim 43, wherein said one or more oxytocin analogue(s) is/are carbetocin is formulated in a powder or aqueous formulation for intranasal delivery.
- 45. (Currently Amended) The method of claim 44, wherein said one or more oxytocin analogue(s) is/are carbetocin is combined in an aqueous formulation with one or more excipients selected from the group consisting of nonoxynol-9, laureth-9, poloxamer-124, octoxynol-9, lauramide DEA, chlorobutanol, glycerol, citric acid, sodium phosphate, methyl paraben, propyl paraben, sorbitol, sodium chloride, and/or sodium acetate for intranasal delivery.
- 46. (Previously Added) The method of claim 44, wherein said carbetocin is formulated with a nonionic surfactant and polysorbate-80 in an aqueous formulation for intranasal delivery.
- 47. (Currently Amended) The method of claim 39, wherein said one or more oxytocin analogue(s) is/are carbetocin is administered in a dose of at least one microgram.
- 48. (Currently Amended) The method of claim 39, wherein said one or more oxytocin analogue(s) is/are carbetocin is administered daily in an intranasal formulation.
- 49. (Previously Added) The method of claim 39, further comprising administering tamoxifen and/or raloxifen to said patient in an amount sufficient to inhibit growth of estrogen-dependent breast cancer in said patient.
- 50. (Currently Amended) The method of claim 49, wherein said one or more oxytocin analogue(s) is/are carbetocin and said tamoxifen and/or raloxifene are administered simultaneously as a mixture